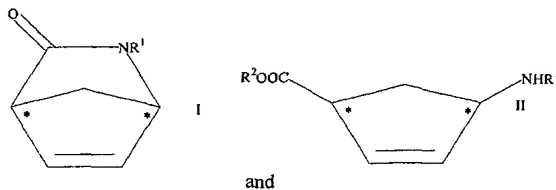
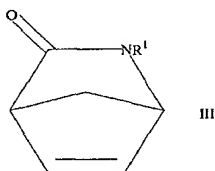


## PATENT

17. (New) A method for forming optically active compounds of the formulae



wherein  $R^1$  is acyl, alkoxycarbonyl or aryloxycarbonyl and  $R^2$  is a  $C_{1-10}$  alkyl, the method comprising treating a racemic lactam of the formula



with a hydrolase and an effective amount of a nucleophile, wherein said nucleophile is a  $C_{1-10}$  alcohol, and a base in a constant pH range to form the optically active compounds of formulae I and II.

18. (New) The method according to Claim 17, wherein a protease or lipase is used as the hydrolase.

19. (New) The method according to Claim 18, wherein a serine protease is used as the protease.

<sup>4</sup>  
20. (New) The method according to Claim <sup>3</sup>19, wherein a subtilisin is used as the serine protease.

<sup>5</sup>  
21. (New) The method according to Claim <sup>1</sup>17, wherein 2-acetyl-2-azabicyclo[2.2.1]hept-5-ene-3-one or 2-ethoxycarbonyl-2-azabicyclo[2.2.1]hept-5-ene-3-one is used as the racemic lactam of formula III.

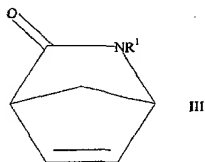
<sup>6</sup>  
22. (New) The method according to Claim <sup>1</sup>17, wherein the treatment of the racemic lactam is conducted in a C<sub>1-10</sub> alcohol or in a mixture of a C<sub>1-10</sub> alcohol with an aprotic solvent.

<sup>7</sup>  
23. (New) The method according to Claim <sup>1</sup>17, wherein the treatment of the racemic lactam is conducted at a temperature of 10 to 60 °C.

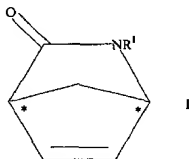
<sup>9</sup>  
24. (New) A method for the formation of optically active 1-amino-4-(hydroxymethyl)-2-cyclopentene derivatives of the formula



wherein R<sup>1</sup> is acyl, alkoxycarbonyl or aryloxycarbonyl, the method comprising treating a racemic lactam of the formula

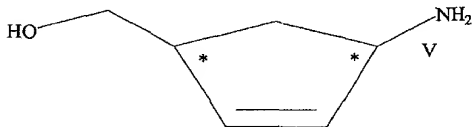


wherein R¹ is acyl, alkoxycarbonyl or aryloxycarbonyl with a hydrolase and an effective amount of a nucleophile, wherein said nucleophile is a C<sub>1-10</sub> alcohol, and a base in a constant pH range to form the compound of the formula



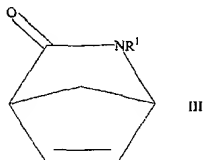
and wherein the compound of formula I is reduced to the compound of formula IV by treatment with a reducing agent.

25. (New) A method for the formation of (1R, 4S)-1-amino-4-(hydroxymethyl)-2-cyclopentene of the formula

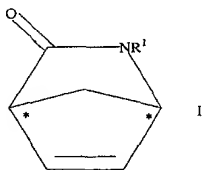


## PATENT

or its salts, the method comprising treating a racemic lactam of the formula



wherein R<sup>1</sup> is acyl, alkoxycarbonyl or aryloxycarbonyl with a hydrolase and an effective amount of a nucleophile and a base in a constant pH range to form the compound of the formula



wherein R<sup>1</sup> is acyl, alkoxycarbonyl or aryloxycarbonyl, wherein the compound of formula I is then reduced to the compound of formula



by treatment with a reducing agent, wherein  $R^1$  is acyl, alkoxycarbonyl or aryloxycarbonyl, and wherein the compound of the formula IV is then hydrolyzed to the compound of formula V.

<sup>8</sup> ~~26~~ 27. (New) The method of Claim <sup>1</sup> ~~17~~, wherein each of the optically active compounds of formulae I and II are isolated after formation.

<sup>10</sup> ~~27~~ 28. (New) The method of Claim <sup>9</sup> ~~24~~, wherein each of the optically active 1-amino-4-(hydroxymethyl)-2-cyclopentene derivatives of formula IV are isolated after formation.

<sup>13</sup> ~~28~~ 29. (New) The method of Claim <sup>12</sup> ~~25~~, wherein the (1R, 4S)-1-amino-4-(hydroxymethyl)-2-cyclopentene of formula V is isolated after formation.

<sup>11</sup> ~~29~~ 30. (New) The method of Claim <sup>9</sup> ~~24~~, wherein the reducing agent is a metal hydride.

<sup>14</sup> ~~30~~ 31. (New) The method of Claim <sup>12</sup> ~~25~~, wherein the reducing agent is a metal hydride.

<sup>15</sup> ~~31~~ 32. (New) A method for forming optically active compounds of the formulae

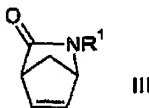


and



wherein  $R^1$  is  $C_{1-4}$  alkanoyl which is substituted with one or more halogen atoms, benzylcarbonyl, phenylcarbonyl, methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl,

butoxycarbonyl or phenyloxycarbonyl and  $R^2$  is a hydrogen atom, the method comprising treating a racemic lactam of the formula



with a hydrolase and an effective amount of water as nucleophile and a base in a constant pH range to form the optically active compounds of the formulae I and II.

1.6 ~~35~~ <sup>31 15</sup> 33. (New) The method according to Claim ~~32~~ <sup>16</sup>, wherein a protease or lipase is used as the hydrolase.

1.7 ~~35~~ <sup>33 17</sup> 34. (New) The method according to Claim ~~33~~ <sup>16</sup>, wherein a serine protease is used as the protease.

1.8 ~~35~~ <sup>33 17</sup> 35. (New) The method according to Claim ~~34~~ <sup>16</sup>, wherein a subtilisin is used as the serine protease.

1.9 ~~35~~ <sup>31 15</sup> 36. (New) The method according to Claim ~~32~~ <sup>16</sup>, wherein 2-ethoxycarbonyl-2-azabicyclo[2.2.1]hept-5-ene-3-one is used as the racemic lactam of formula III.

2.0 ~~35~~ <sup>31 15</sup> 37. (New) The method according to Claim ~~32~~ <sup>16</sup>, wherein treatment of the racemic lactam is conducted in water, a buffer solution or in a mixture of these with an aprotic solvent.